(12) UK Patent Application (19) GB (11) 2 217 630(13)A

(43) Date of A publication 01.11.1989

- (21) Application No 8810085.4
- (22) Date of filing 28.04.1988
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- (51) INT CL4 B05D 3/00
- (52) UK CL (Edition J) B2E EFA E1201 E1217 E1311 E1327 E1329 E1701 E436T E443S E450T E468T E474S E489T E491T U1S S1129
- (56) Documents cited None
- (58) Field of search UK CL (Edition J) B2E INT CL BOSD

(54) Microsphere coating material

(57) A method for the manufacture of materials for shoe linings and counter linings comprises applying to a fibrous substrate a coating composition comprising a dispersion of a film-forming polymer, in which are dispersed microspheres which are expansible on heating, subjecting the material in a first heating stage to a temperature at which the film-forming polymer forms a cohesive film but at which the microspheres do not expand, and then subjecting the material in a second heating stage to a temperature at which the microspheres expand and a microcellular structure is developed.

The shoe linings and counter linings manufactured by this method are particularly suitable fo the manufacture of linings with a coloured surface which may be sueded.

The present invention relates to improvements in or relating to shoes and the manufacture thereof and is especially concerned with materials used in the manufacture of shoes which are subject in wear to abrasion, for example shoe linings, in particular shoe counter-linings. "shoe" is used herein to denote outer footwear generally 10 whether ready for wear or in the course of manufacture.

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Many shoes have uppers comprising a outer and a In ladies' court shoes, whether the uppers are lining. lined all over the inside or not, the counter region often lining of suitable material has known 15 counter-lining. A counter-lining is preferably made of a material having a high coefficient of friction with respect to the foot and hose of a wearer to assist in retention of the shoe on the foot in wear; counter-lining materials preferably have a good resistance to abrasion and an 20 appearance pleasing to the eye of a potential purchaser of a shoe in which the counter-lining has been incorporated.

A further requirement which has arisen relatively recently, in particular as a result of current fashions, is materials for shoe linings, in particular 25 counter-linings, to have a coloured surface as hereinafter defined, and it may be desired that this coloured surface should be combined with a sueded surface.

Prior art linings are known which may be produced with a sueded surface and, of these, the lining materials 30 which are manufactured by impregnating a non-woven fibrous material with an aqueous dispersion comprising carboxylated rubber, a curing system for the rubber and a vinyl chloride polymer resin, drying the impregnated material and curing the carboxylated rubber, followed where 35 required by sueding off of the surface, have proved to be particularly successful.

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The materials which are manufactured by this method are particularly suitable for the production of shoe linings and counter linings in pastel colours. The base materials are formed from white or light coloured fibres, polypropylene/polyester typically polymers, inevitably pastellise the product, and this effect becomes particularly marked when the surface is sueded.

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An object of the present invention is to provide a method for the manufacture of materials for shoe linings 10 and counter linings, which may optionally have a coloured surface and which may optionally be produced with a sueded surface while retaining the coloured surface where this is provided.

The present invention provides a method for the 15 manufacture of materials for shoe linings and counter linings which method comprises applying to a substrate a coating composition comprising a dispersion of a film-forming polymer, in which are dispersed microspheres which are expansible on heating, subjecting the material in 20 a first heating stage to a temperature at which the film-forming polymer forms a cohesive film but at which the microspheres do not expand, and then subjecting the material in a second heating stage to a temperature at which the microspheres expand and a microcellular structure is developed.

The present invention further provides materials for shoe linings and counter linings, which materials are manufactured by a method which comprises applying to a substrate coating composition comprising fibrous а 30 dispersion of a film-forming polymer, in which dispersed microspheres which are expansible on heating, subjecting the material in a first heating stage to a temperature at which the film-forming polymer forms a cohesive film but at which the microspheres do not expand, and then subjecting the material in a second heating stage to a temperature at which the microspheres expand and a microcellular structure is developed.

The method according to the invention is preferably a method for the manufacture of materials for shoe linings and counter linings which have a coloured surface as herein defined, in which the coating composition applied to the substrate further comprises a pigment.

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The materials for shoe linings and counter linings manufactured by the method according to the 10 invention preferably have a coloured surface as herein defined, and are manufactured by a method which comprises applying to the fibrous substrate a coating composition which further comprises a pigment.

By the term "coloured surface" as used herein, is
15 meant a surface which is pigmented to give a colour which
is not pastellised by the effect of white or light-coloured
fibres which are present in the fibrous substrate and which
affect the appearance of the surface of the lining.

The fibrous substrate to be used in the method 20 according to the invention preferably comprises a bonded non-woven or spun bonded fabric. Suitable materials for the fibres include polypropylene/polyester polymers.

The film-forming component applied to the substrate preferably comprises an aqueous dispersion of a 25 film-forming polymer. Suitable dispersions include latices and aqueous polyurethane dispersions.

Where the film-forming component is applied in the form of an aqueous dispersion, the first heating stage is a drying stage and is suitably carried out at a temperature below 100°C, preferably below 90°C at which temperature the microspheres are not expanded. Expansion of the wet coating should be avoided, as the final coating is more uniform if the coating is dried first. The dried and unexpanded coating is subsequently heated in the second heating stage to a temperature at which the microspheres

are expanded, preferably above 120°C, more preferably 140° to 150°C.

Where the material is required to have a sueded surface, the expanded coating is sueded, for example with emery paper, to produce a level surface, with exposed fine cell structure, which surface has a uniform, matt and intense colour.

Suitable methods by which the composition used according to the invention can be coated onto the substrate include, in particular, knife coating, for example with a knife on a roll, or rotary screen printing. If rotary screen printing is used, it is helpful to position a smoothing blade after the screen head to remove any pattern left by the screen.

During the expansion stage of the method according to the invention, that is the second heating stage, there is a possibility that the base material will curl along its length, so that stentering is necessary. This tendency for curling, and the consequent need for stentering, is removed if the expansion stage is carried out on a drum, for example a Fleissner drum.

The invention will now be further described with reference to the following specific examples:-

Example 1

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A coating system having the following composition was stirred and then milled in a colloid mill to disperse the microspheres:

		Parts by Weight	
		<u>Wet</u>	Dry
30	Nitrile Latex 8250	14,000	5,600
	PVC Latex 7151	6,000	3,540
	Black Pigment Dispersion	800	200
	30% Expancel 461 WU	3,066	919
	10% Viscalex HV30	1,470	147
35	DB 110 (50%) Antifoam	25.3	12.6

Nitrile Latex 8250 is a cross-linkable carboxylated nitrile rubber supplied by Bayer A.G.

PVC Latex 7151 is a polyvinyl chloride latex supplied by Polysar.

Expancel 461 WU is an expansible microsphere material in wet unexpanded form, supplied by Nobel Industries, Sweden.

Viscalex HV30 is a polyacrylic thickening agent supplied by Allied Celloids Ltd.

DB 110 is a silicone antifoam agent supplied by Dow Corning.

Immediately before use, the following components were stirred into the coating system:

		Parts by Weight
15	0.5% .880 Ammonia	127
	5.9% Beetle Resin 338	1,495
	5% Viscalex HV30 (10%)	1,270

Beetle Resin 338 is a melamine formaldehyde 20 condensate supplied by BIP Chemicals, and was used to cross-link the carboxylated nitrile rubber.

The ammonia was added to solubise the HV30 polyacrylic acid thickening agent.

The viscosity of the latex/Expancel blend after the final additions and stirring was 25,000 to 30,000cps when measured on a Brookfield viscometer, spindle 6, speed 10.

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The coating blend was coated onto a bonded non-woven fabric substrate, 0.5mm thick, using a Meyer bar.

The substrate was coated at a coating weight of 140 to 170gm per square metre and dried at 90°C. At this temperature, the coating dried without expansion of the microspheres. After drying, the thickness of the coated material was from 0.60 to 0.65mm.

35 The coated substrate was then heated at 140° to 150°C for 1 to 2 minutes, to expand the microspheres. The

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gauge increased to 1.1 to 1.2mm. The expanded material was then heated for a further 5 minutes, to cure the nitrile rubber and develop optimum abrasion resistance. The material was finally sueded to a gauge of 0.8mm.

The resultant material had intense surface colouration, uniform thickness and a sueded surface, with good abrasion resistance.

Example 2

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A coating system having the following composition 10 was stirred and then milled in a colloid mill to disperse the microspheres:

		Parts by Weight	
		Wet	Dry
	Nitrile Latex 2890	250	100
15	30% Expancel 461 WU	33.3	10
	Black Pigment Dispersion	10	2.5
	10% Viscalex HV 30	50	5
	DB 110 (5%) Antifoam	0.5	0.03

20 Just prior to coating, the following curing ingredients were added:

		Parts	by Weight
		Wet	Dry
	0.88 Ammonia (as required)		
25	50% Sulphur Dispersion 2012	4	2
	50% Zinc Oxide Dispersion 209	7 6	3
	50% ZDC Accelerator		
	Dispersion 12	2	1
	50% ZMBT Accelerator 1831	2	1
30	Beetle Resin 338	18.75	14.1
	10% Viscalex HV 30	69	6.9

The Nitrile Latex 2890 is supplied by Bayer A.G. and the Sulphur 2012, Zinc Oxide 2097, ZDC 12, and ZMBT 1831 are supplied by Rubber Latex Ltd.

The viscosity of the latex/Expancel blend after the final additions and stirring was 25,000 to 30,000cps. when measured on a Brookfield viscometer, spindle 6, speed 10.

Coating, drying and heating were then carried out as described in Example 1. The gauge increased to 1.1 to 1.2mm. Final curing was carried out as in Example 1. The material was finally sueded to a gauge of 0.8mm.

The resultant material had intense surface 10 colouration, uniform thickness and a sueded surface with good abrasion resistance.

Example 3

The procedure of Example 2 was repeated, with the variation that the 100 parts by weight dry of Nitrile 15 Latex 8250 was replaced by 75 parts by weight dry of Nitrile Latex 2890 and 25 parts by weight of Nitrile Latex 8250.

As compared to the coating produced according to Example 2, this material had slightly increased surface 20 coating hardness.

Example 4

The procedure of Example 1 was repeated, with the variation that the Nitrile Latex used in Example 1 was replaced by 5600 parts by weight dry of Doverstrand Revinex 5043, a styrene - butadiene - acrylonitrile prepolymer latex.

The resultant material had intense surface colouration, uniform thickness and a sueded surface with good abrasion resistance.

30 Example 5

A coating system having the following composition was stirred and then milled in a colloid mill to disperse the microspheres:

		Parts b	y Weight
		Wet	Dry
	Permatex EX4844 PU dispersion	250	87.5
	Permatex UA9048 Crosslinker	7.5	5.25
5	30% 551 Expancel	33.3	10
	Coloured pigment dispersion	10	2.5
	PB110(5%) Antifoam	0.5	0.03
	0.88 Ammonia (as required)		
	10% Viscalex HV30	50	5

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Permatex EX4844 and UA9048 are supplied by Stahl Chemicals Ltd.

Coating was carried out as described in Example 1, and the coated substrate was dried at 60 or 70°C.

The coated substrate was then blown at 120°C, at which temperature the microspheres were expanded and the polyurethane completely cured.

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CLAIMS:-

- A method for the manufacture of materials 1. for shoe linings and counter linings, which method comprises applying to a fibrous substrate a coating composition comprising a dispersion of a film-forming polymer, in which are dispersed microspheres which are expansible on heating, subjecting the material in a first heating stage to a temperature at which the film-forming 10 polymer forms a cohesive film but at which the microspheres do not expand, and then subjecting the material in a second heating stage to a temperature at which the microspheres expand and a microcellular structure is developed.
- 15 2. A method as claimed in Claim 1, in that the substrate comprises a bonded non-woven or spun bonded fabric.
- 3. A method as claimed in Claim 1 or Claim 2 in 20 which the coating composition further comprises a pigment.
 - A method as claimed in any of Claims 1 to 3 4. which further comprises the step of sueding the surface.
- 25 5. A method as claimed in any of Claims 1 to 4 in which the microspheres are dispersed in an aqueous dispersion.
- A method as claimed in any of Claims 1 to 5 6. 30 in which the microspheres are dispersed in a latex or an aqueous polyurethane dispersion.
- A method as claimed in any of Claims 1 to 6 in which the material is heated in the first heating stage 35 to a temperature below 100°C.

8. A method as claimed in Claim 7 in which the material is heated in the first heating stage to a temperature below 90°C.

9. A method as claimed in any of Claims 1 to 8 in which the material is heated in the second heating stage to a temperature above 120°C.

10. A method as claimed in Claim 9 in which the 10 material is heated in the second heating stage to a temperature of from 140 to 150°C.

11. A material for shoe linings and counter linings which has been manufactured by a method as claimed in any of Claims 1 to 10.

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